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**Reading targeted DNA damage in the active demethylation
pathway: role of accessory domains of eukaryotic AP endonucleases
and thymine-DNA glycosylases**

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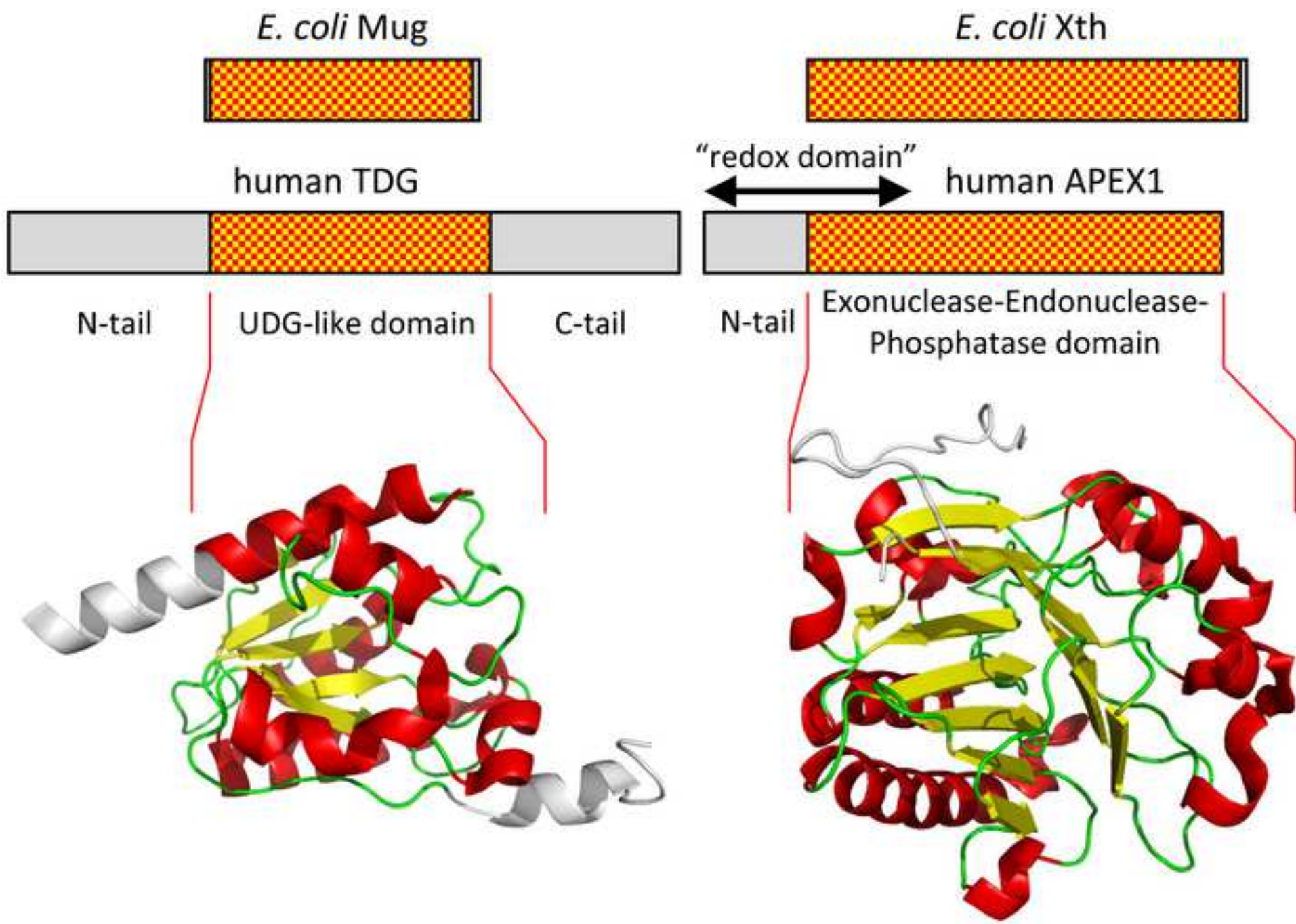
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disorder

ABBREVIATIONS

ROS, reactive oxygen species; AP, apurinic/aprimidinic site; BER, base excision repair;
NIR, nucleotide incision repair; APE1 (a.k.a. APEX1, HAP-1, or Ref-1), human major AP
endonuclease 1; TDG, T:G mismatch-specific thymine-DNA glycosylase. mC, 5-
methylcytosine; hmC, 5-hydroxymethylcytosine; fC, 5-formylcytosine; caC, 5-carboxylcytosine;
fU, 5-formyluracil; hmU, 5-hydroxymethyluracil; TF, transcription factor; CGIs, CpG islands;
Rrp1, *Drosophila* AP endonuclease 1.



Highlights_Popov et al 2019

HIGHLIGHTS

- Eukaryotic BER proteins contain extended non-catalytic tails absent from bacterial and archaeal homologs;
- The tails of TDG and APE1 homologs are more diverse than their catalytic cores and contain diverged short repeats;
- The recombination function cannot be universally ascribed to the N-tail of insect APE1 homologs;
- The tails of TDG and APE1 participate in the formation of oligomeric aggregates on DNA and enable active epigenetic demethylation;
- The APE1 and TDG aggregates on DNA might be involved in liquid phase separation.

Abstract

1 Base excision DNA repair (BER) is an important process used by all living organisms to
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4 remove non-bulky lesions from DNA. BER is usually initiated by DNA glycosylases that excise
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6 a damaged base leaving an apurinic/apyrimidinic (AP) site, an AP endonuclease then cuts DNA
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8 at the AP site, and the repair is completed by correct nucleotide insertion, end processing, and
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10 nick ligation. It has emerged recently that the BER machinery, in addition to genome protection,
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12 is crucial for active epigenetic demethylation in vertebrates. This pathway is initiated by TET
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14 dioxygenases that oxidize the regulatory 5-methylcytosine, and the oxidation products are treated
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16 as substrates for BER. T:G mismatch-specific thymine-DNA glycosylase (TDG) and AP
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18 endonuclease 1 (APE1) catalyze the first two steps in BER-dependent active demethylation. In
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20 addition to the well-structured catalytic domains, these enzymes possess long tails that are
21
22 structurally uncharacterized but involved in multiple interactions and regulatory functions. In
23
24 this review, we describe known roles of the tails in TDG and APE1, discuss the importance of
25
26 order and disorder in their structure and consider the evolutionary aspects of these accessory
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28 protein regions. We also propose that the tails may be important for the enzymes'
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30 oligomerization on DNA, an aspect of their function that only recently gained attention.
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1. Introduction

DNA repair, a system of several partially overlapping enzymatic pathways, has evolved to protect genetic material from ongoing damage and is present in some form in all cellular organisms [1]. Of these pathways, base excision repair (BER) removes small non-bulky lesions from DNA; lesions of this kind are the most abundant source of mutations and are mostly unavoidable since they are generated through nucleotide hydrolysis and oxidation [1-4]. In the simplest form of BER, the damaged base is located and excised by one of the lesion-specific enzymes belonging to the group of DNA glycosylases, the nascent abasic (apurinic/apyrimidinic, AP) site is cut by an AP endonuclease, the correct dNMP is inserted by a DNA polymerase with simultaneous removal of the hanging deoxyribose remnant, and the resulting nick is sealed by a DNA ligase [1-4]. Several variations on this general mechanism operate in mammalian cells, differing in the requirements for DNA glycosylases and AP endonucleases and generating single-strand breaks with different termini [2-5]. Deficiencies in the key components of BER are embryonic lethal [6-7] while certain functional variants are associated with an increased risk of cancer [8-10].

In recent years, much attention has been drawn to non-canonical functions of BER, not related to genome integrity maintenance in a strict sense. Obviously, the versatile BER machinery can be easily adopted to processing regulated targeted DNA modification in the cell. BER has been shown to be involved in somatic hypermutation and class switch recombination during immunoglobulin gene maturation [11-12], virus life cycle and cell antiviral defense [13-14] and active epigenetic cytosine demethylation [15-17]. In higher eukaryotes, this last pathway is initiated when the regulatory DNA base 5-methylcytosine is oxidized by TET family dioxygenases and/or deaminated by AID and APOBEC family deaminases and then removed preferentially by mismatched thymine-DNA glycosylase, TDG. The resulting AP site is then

1 hydrolyzed by the major AP endonuclease APE1 and further processed *via* regular BER
2 mechanisms.

3 Many protein molecules in living organisms consist of two or more structural domains,
4 tens to hundreds amino acids long [18-19]. A domain is usually defined as a separate protein part
5 that has a defined structure and function and may evolve largely independently of the rest of the
6 protein. In the case of enzymes, their catalytic activity is often localized in one or several
7 domains, which form a core fragment, while other domains of the same protein may participate
8 in activity regulation, protein–protein interactions, localization in the cell, etc. In multidomain
9 proteins, often the function of the core fragment is known whereas the roles of other domains are
10 unclear. The development of methods for protein domain prediction based on homology and
11 structural modeling [20-21] led to description of hundreds of domains of unknown functions.
12 Moreover, there are many cases when a protein possesses known domains and long tails or
13 linkers that are not identified as domains. Structurally they may be true domains (i. e., have a
14 defined structure and be separated from other domains), but more often they are disordered and
15 either not solved by X-ray crystallography or prevent crystallization and thus have to be removed
16 to solve the core structure of the protein. The lack of structural information much complicates
17 the analysis of their functions.

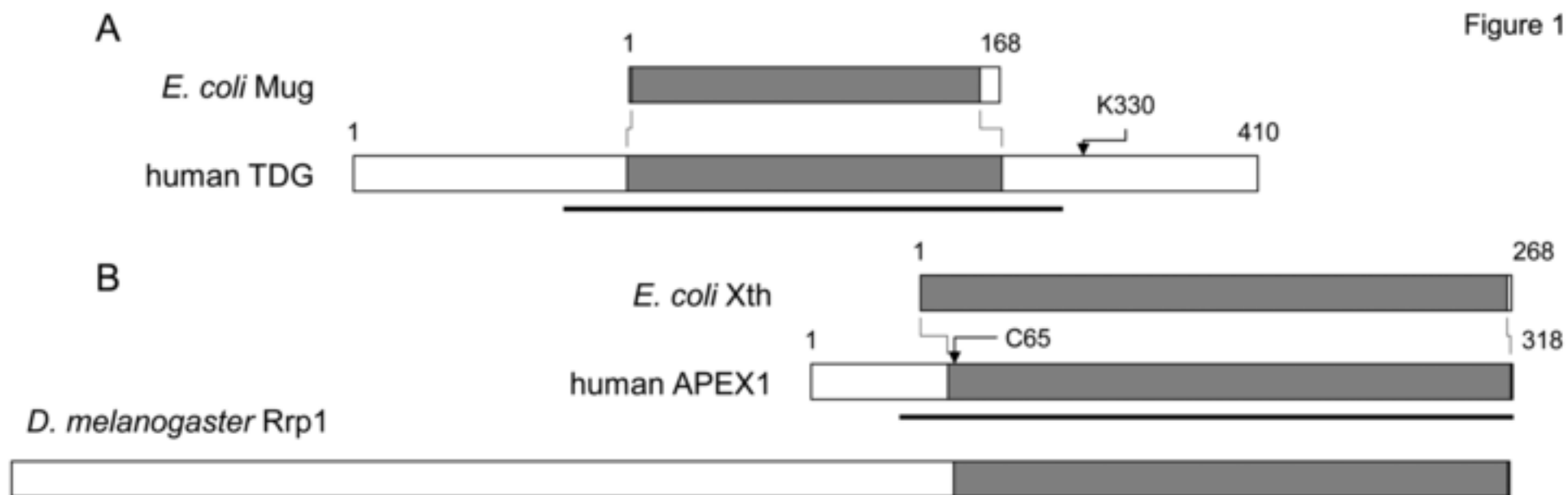
18 Available structural data show that many eukaryotic BER enzymes consist of a well-
19 resolved core domain (or several domains) responsible for catalysis and/or DNA binding, and
20 terminal tails that are, in most cases, disordered [22-23]. These tails generally can be removed
21 without affecting the activity of the enzyme, and the bacterial counterparts of eukaryotic BER
22 enzymes are most often compact proteins with little extra sequence outside the catalytic domain
23 borders. The functions of these tails in eukaryotic BER, except in a very few occurrences, remain
24 enigmatic but they are usually considered to be involved either in protein–protein interactions
25 [24-25] or in non-specific DNA binding contributing to the affinity of the protein for DNA or
26 facilitating its movement along DNA [26-27].

1 Based on the structure of the core domain, human DNA glycosylases can be assigned to
2 four superfamilies: UDG-like (uracil-DNA glycosylase homologs), HhH (containing a helix-
3 hairpin-helix DNA binding motif), H2tH (containing a helix-two-turn-helix DNA binding
4 motif), and FMT_C (homologs of the C-terminal domain of methionyl-tRNA formyltransferase).
5
6 AP endonucleases belong to two major superfamilies, EEP (exonuclease-endonuclease-
7 phosphatase) and TIM (triosephosphate isomerase) barrel. Human TDG and APE1, which
8 initiate the BER stage of active DNA demethylation, belong to the UDG-like and EEP
9 superfamilies, respectively. Both TDG and APE1 fit the structural paradigm of the ordered core
10 domain and disordered tails (Figure 1). In this review, we summarize what is known about the
11 functions of the tails of TDG and APE1 and discuss their evolution and the possible role in the
12 regulation of epigenetic demethylation¹.
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28 **2. TDG: a hub of epigenetic DNA demethylation**

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31 Methylation of cytosine at the C5 position with the formation of 5-methylcytosine (mC)
32 in DNA provides the molecular basis of epigenetic regulation of gene expression in several
33 major groups of eukaryotes [28-30]. DNA methylation is essential for the organism
34 development, cell differentiation, genomic imprinting and suppression of mobile genetic
35 elements. A drawback of this mode of regulation is that spontaneous deamination of mC
36 generates thymine, resulting in a T:G mismatch which, if not repaired, leads to C→T transitions
37 at CpG dinucleotides [31]. In mammalian cells, two DNA glycosylases, mismatch-specific
38 thymine-DNA glycosylase (TDG) and methyl-binding domain protein 4 (MBD4/MED1) prevent
39 the mutagenic impact of mC deamination by excising T from T:G mispairs in CpG context thus
40 initiating BER that restores C [32-33].
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58 ¹ In the literature, the term “domain” is often used indiscriminately to describe true structural domains, or
59 structurally non-discrete parts of the protein responsible for specific functions, or disordered regions. Here, we will
60 consistently refer to structural domains as “domains” (or “cores” specifically when describing the main catalytic
61 domains), parts responsible for specific functions as “parts”, and terminal disordered regions as “tails”.

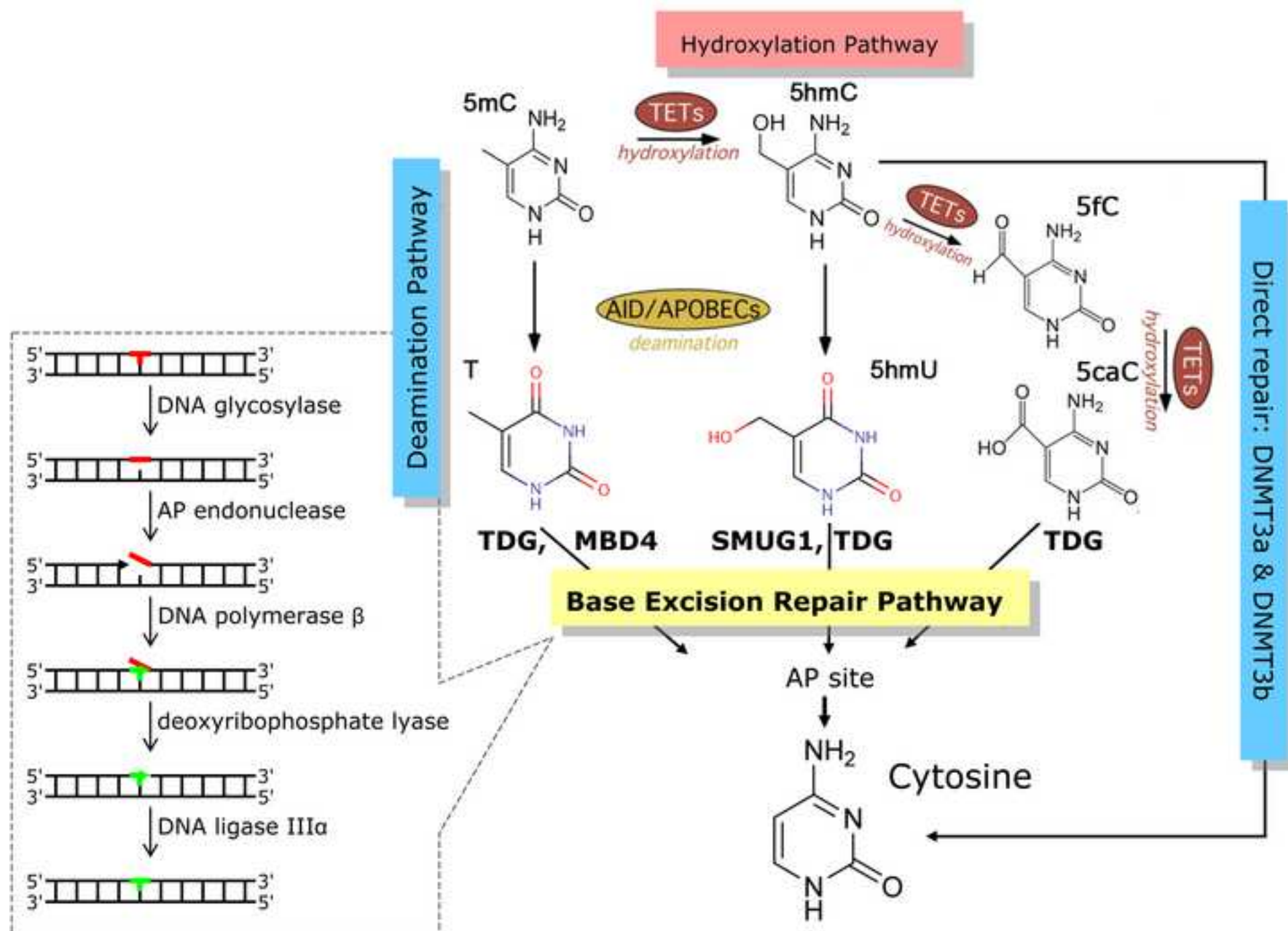


1 enzymatic activity excising Thy bases from T:G mispairs [32,34-35], purified and cloned,
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3 revealing significant homology with *E. coli* mismatched uracil–DNA glycosylase (Mug) [36].
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5 Subsequent discovery of additional Mug/TDG homologs showed that they form a separate
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7 family within the uracil–DNA glycosylase-like superfamily [37-39]. A more detailed enzyme
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9 activity inquiries showed that TDG exhibits wide substrate specificity, excising 3,*N*⁴-
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11 ethenocytosine [40-41], thymine glycol [42], 5-hydroxycytosine [43], 7,8-dihydro-8-oxoadenine
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13 [44], mismatched U [35] and its C5-modified derivatives [43,45]. In contrast, MBD4 has narrow
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15 DNA substrate specificity, excising T, U, 5-fluorouracil and 5-hydroxymethyluracil in a XpG
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17 context from duplex DNA [33,46-47].
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23 Until recently, TDG had been considered as functional “back-up” for the major vertebrate
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25 uracil–DNA glycosylase, UNG, with somewhat broadened damaged base specificity. The
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27 discovery of the central place of TDG in active epigenetic demethylation revealed what is likely
28
29 the true biological function of at least the vertebrate homologs; no similar role outside the basic
30
31 BER is presently known for bacterial Mug proteins. Active DNA demethylation in vertebrates is
32
33 dependent on the Ten-eleven translocation family of proteins (TETs; TET1, TET2, and TET3),
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35 iron/ α -ketoglutarate-dependent dioxygenases that convert mC to 5-hydroxymethylcytosine
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37 (hmC) and then further oxidize it to 5-formylcytosine (fC) and 5-carboxylcytosine (caC) [48-51]
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39 (Figure 2). TDG excises fC and caC from CpG context with high efficiency, suggesting direct
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41 involvement of the TDG-initiated BER in the active erasure of mC epigenetic marks [50,52-57].
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43 In addition, mC and its C5-oxidized derivatives can be deaminated by enzymes belonging to the
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45 activation-induced cytidine deaminase (AID) / Apolipoprotein B mRNA-editing enzyme
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47 complex (APOBEC) family of cytidine deaminases [58-60], with at least 5-hydroxymethyluracil
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49 (hmU) and 5-formyluracil (fU) also being substrates for TDG [45,53-54,61-62]. In agreement
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51 with these observations, *Tdg* knockouts in mice are embryonic lethal due to aberrant *de novo*
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53 DNA methylation of CpG islands in promoters of developmental genes and failure to establish
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Figure 2

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and/or maintain cell type-specific gene expression programs during embryonic development [62-63].

In addition to TETs/TDG-dependent active erasure of 5mC residues, a second mechanism was shown, in which AID catalyses deamination of 5mC to thymine generating a G•T mismatch base pair that is repaired by MBD4 [58]. Moreover, it was shown that AID/APOBEC catalyze deamination of 5hmC to hmU residue, which is in turn excised by the MBD4, and single-strand specific monofunctional uracil-DNA glycosylase 1 (SMUG1) [59,62]. MBD4/MED1 is a bipartite protein that belongs to the family of methyl-CpG-binding domain (MBD) proteins, and consists of an N-terminal MBD domain that is linked to a C-terminal DNA glycosylase domain [33,46]. Mutations of MBD4 gene were identified in tumors with defective DNA mismatch repair, however, knockout of MBD4 in mouse causes only a small 2-3 fold increase in C→T mutations at CpG sites and did not increase mini-satellite instability implying that MBD4 may act as a modifier and not as driver of tumorigenesis [64-65]. MBD4 interacts with the mismatch repair protein MLH1 [66] and DNA methyltransferases DNMT1 and DNMT3b [67-68] suggesting a potential link between post-replication repair and DNA methylation. Noteworthy, the interaction between MBD4 and DNMT1 is involved in the repair of oxidative DNA damage and control of gene expression at methylated CpG islands [69]. These observations suggest that MBD4 is a back up DNA glycosylase involved in the epigenetic regulation *via* removal of oxidation and deamination products of 5mC.

Structurally, TDG consists of the Mug-like catalytic core 171 amino acid residues (aa) long and two tails, N-terminal (N-tail, 123 aa) and C-terminal (C-tail, 116 aa). The available crystal structures of TDG leave the tails mostly unresolved ([Figure 1](#)) [54-56,70-76]. NMR studies suggest that the N-tail is largely disordered, however, its part proximal to the catalytic core shows interactions both with the core and with bound DNA [75,77]. No experimental information regarding the C-tail is available.

1 From the moment of TDG discovery, the N-tail has been suspected to be somehow
2 involved in the substrate specificity: progressive deletions into the N-tail eventually abolish the
3 activity on T:G mismatches but hardly affect or even modestly increase the uracil excision from
4 U:G mismatches [37]. Residues 82–110 appear to be critical for the efficient T:G binding by
5 TDG [75]. Additionally, substrate processing by TDG can be regulated by the N-tail at the level
6 of enzyme turnover: full-length TDG has very high affinity for the AP site left after base
7 excision and normally requires stimulation by APE1 for product release [78-79], whereas
8 deletion of the N-tail dramatically increases the turnover on the U:G substrate [80]. The
9 importance of the N-tail for the processing of substrates other than T:G and U:G remains largely
10 unknown.
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22 Another factor facilitating TDG enzymatic turnover is its covalent modification by small
23 ubiquitin-like modifier proteins SUMO-1 and SUMO-2/3 at Lys330 in the C-tail, a residue
24 absolutely conserved within Deuterostomia [80-81]. The sumoylation drastically reduces TDG
25 affinity for AP sites or normal and damaged DNA and modifies its substrate specificity,
26 enhancing the cleavage of U:G and suppressing the cleavage of T:G mismatches [81]. It also
27 improves TDG stimulation by APE1 [81]. Weakening a network of hydrogen bonds between
28 SUMO and the C-tail by site-directed mutagenesis abolishes the effect on sumoylation on DNA
29 binding [71]. A comparison of the structures of unmodified and sumoylated TDG [71,74-75]
30 suggest that the rigidly attached SUMO molecule clashes with DNA near the lesion, which likely
31 should lead to the protein–DNA complex falling apart. Noteworthy, the sumoylation effect on
32 TDG turnover also requires an intact N-tail [80,82], and a site for non-covalent SUMO binding is
33 found in the N-terminal part of the catalytic core [82]. As evidenced by NMR, covalent
34 conjugation and non-covalent binding of SUMO to the respective sites induce distinct
35 conformational changes in the TDG tails [83]. During active demethylation, SUMO1 is first
36 attached to XRCC1, the main BER scaffold protein, which enhances the assembly of the BER
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complex involving TDG, to which SUMO1 is later transferred to promote further repair steps [84].

Disordered tails in proteins are often involved in protein–protein interactions. TDG is implicated in the regulation of transcription by structurally diverse factors such as retinoic acid receptors (RAR α /RXR α) [85], estrogen receptor ER α [86], several other nuclear receptor family proteins [87], basic leucine zipper protein c-Jun [85,88], homeodomain-containing thyroid transcription factor 1 [89], p53 and p73 [90]. The interactions with the individual transcription factors have been mapped to the catalytic core. However, TDG complexes with transcription factors can be elaborated by including transcriptional co-activators – CBP/p300 lysine acetylases that initiate chromatin remodeling. Taken alone, TBP can bind CBP/p300 through both N- and C-tails [91]. The resulting complex is competent in both BER and histone acetylation. TDG enhances the CBP/p300 transcriptional activity and is in turn acetylated by CBP/p300, negatively regulating the recruitment of APE1. The TDG–CBP/p300 complexes can also include RAR α /RXR α [92] and LEF1/TCF4, a complex involved in the Wnt/ β -catenin signaling pathway [93]. Although the role of the glycosylase activity is not evident in all these cases, multiple interactions of TDG with transcription regulation machinery including CBP/p300 may couple DNA demethylation with chromatin unfolding for gene activation.

Finally, TDG expression is strictly cell-cycle regulated: it is present in cells throughout the G2-M and G1 phases, but rapidly disappears in the S phase [94]. This regulation involves TDG proteasomal degradation initiated by the association of TDG with PCNA through the “PIP degron”, a PCNA-interacting peptide that triggers recognition by CRL4^{Cdt2} E3 ubiquitin ligase upon binding [95-96].

CpG islands (CGIs) are short interspersed CG-rich genomic regions that are abundant near the transcription start sites of genes [97]. In mammalian genomes, CGIs are typically 500–3,000 base pairs in length, and have been found in or near half of the promoters [98]. As key elements in transcription regulation, CGIs are enriched in permissive histone modifications, poor

1 in DNA cytosine methylation and contain multiple sites for transcription factors [99]. Despite
2 CpG are significantly underrepresented in mammalian genomes, their enrichment at transcription
3 start sites appeared early in the evolution of vertebrates [100]. It was hypothesized that the
4 emergence and stabilization of CGIs could be due to a hypodeamination regime associated with
5 low level of DNA methylation or GC-biased gene conversion – a non-reciprocal copying of a
6 DNA sequence from one homologous chromosome to the other during meiotic recombination
7 [101-102]. At present, the molecular mechanisms underlying the CpG enrichment at regulatory
8 regions and the emergence of CGIs in evolution remain unclear. Interestingly, the nuclear
9 fraction of TDG associates tightly with euchromatin and binds to CpG-rich promoters of
10 transcribed genes including the pluripotency and developmental genes [63,91]. *Tdg*^{-/-} mouse
11 embryonic fibroblasts show imbalanced histone modifications and CpG methylation at the
12 promoters of the affected genes, suggesting that the TDG-initiated BER is required to support
13 correct methylation patterns and chromatin remodeling in specific DNA regions.
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30 Recently, it has been demonstrated that TDG, and to a lesser extent MBD4, is able to
31 excise T opposite to various adenine lesions with good efficiency from the specific 5'-TpG-3'/5'-
32 CpX-3' context (where X is hypoxanthine (Hx), ethenoadenine, 8-oxoadenine, or AP site) [103].
33 This ability was classified as aberrant DNA repair, since it can lead to misincorporation opposite
34 the damaged base [104]. The isolated catalytic domain of TDG (residues 111–308) lacking both
35 N- and C-tails, while maintaining the uracil removal activity, excises T from these substrates
36 very weakly, suggesting that the N- and C-terminal portions of TDG are required for the aberrant
37 activity. In vitro reconstitution of TDG-initiated BER with DNA containing a T:Hx pair results
38 in the misincorporation of C opposite to Hx. Furthermore, comparison of the spontaneous
39 genome-wide mutation spectra with those at the CGIs revealed that CGIs exhibit a strong
40 mutational bias for TpG, CpA→CpG mutations [103]. These findings demonstrate that TDG
41 may catalyze sequence context-dependent aberrant removal of T that results in the appearance of
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1 a CpG dinucleotide, thus providing a plausible mechanism for the evolutionary origin of CpG
2 islands in mammalian genomes.
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6 **3. APE1: a dual function enzyme**

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8 AP endonuclease 1 (variously called APE1, APEX1, HAP-1 or Ref-1 in the literature) is
9 the major AP endonuclease supporting BER in vertebrate cells both for genome defense and for
10 active DNA demethylation, and is also involved in RNA biogenesis [105-106]. APE1 was
11 independently discovered as an AP site-specific DNA endonuclease homologous to the *E. coli*
12 Xth protein [107-108] and as a redox factor (Ref-1) that regulates the DNA-binding domains of
13 the Fos and Jun subunits of AP-1 transcription factor [109]. Based on the biochemical
14 characterization of N- and C-terminal deletion mutants, the APE1 polypeptide can be subdivided
15 into two functionally distinct and partially overlapping regions, with the N-terminal part
16 (residues 1–127) possessing the “redox activity”, and the C-terminal part (residues 61–318)
17 responsible for the AP site cleavage activity [110-111]. Structurally, however, the redox part
18 does not fold into a separate domain, and APE1 contains an Xth-like catalytic core (residues 62–
19 316) and an N-tail unresolved in the available crystal structures over the first 2/3 of its length
20 [112-123].
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40 The redox function localized to the N-tail and the adjacent part of the catalytic domain
41 has been a matter of extensive studies, which nevertheless still have not come to an agreeable
42 model; even the redox nature of this activity is controversial. Initially, it was shown that spurious
43 oxidation of Cys272 in Jun and Cys154 in Fos subunits of AP-1 transcription factor prevent its
44 binding to DNA [124], and that this binding can be restored by Ref-1, a protein factor that turned
45 out to be identical to APE1 [109,125]. Truncated APE1 containing only residues 1–127 is fully
46 able to stimulate AP-1 DNA binding [110]. The N-tail is essential for the stimulation of AP-1
47 binding to DNA, and its deletion results in a > 10-fold loss in the stimulation activity but does
48 not affect AP site cleavage [110-111,126]. In addition to AP-1, APE1 has been reported to
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1 enhance DNA binding by NF- κ B [109,127-129], p53 [130-132], Egr-1 [133], Pax-5, Pax-8 [134-
2 135], HIF-1 α and HLF [136-137], and YB-1 [138], although the localization of the stimulation
3 activity to the Ref-1 part in many of these cases has not been confirmed.
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6 Based on mutagenesis results, APE1 was initially proposed to activate AP-1 and NF- κ B
7 via a redox mechanism by mediating the reduction of a Cys residue in their DNA-binding
8 domains [115,126]. A disulfide bond would form in APE1 between Cys65 (residing in the
9 catalytic domain) and presumably Cys93, which is then regenerated by thioredoxin as a reducing
10 molecule [139-140]. However convenient, this model is plagued by a number of contradictory
11 findings. First, crystallographic studies revealed that Cys65 and Cys93 are buried inside the
12 APE1 globule and are unlikely to interact with other proteins [112-113]. Second, the ablation of
13 either the homologous Cys64 residue in mouse Apex1, or of all seven cysteines in human APE1
14 does not affect the stimulation of DNA binding by AP-1 and NF- κ B [129,141]. Third, the
15 *Arabidopsis* AP endonuclease ARP, which is only distantly related to APE1, lacks a Cys65
16 homolog and shares very little homology with the Ref-1 part of APE1, is able to stimulate the
17 DNA-binding activity of human AP-1 [142]. Finally, at least in AP-1, the Cys oxidation does not
18 produce a disulfide bond but rather involves other oxidized thiol derivatives [124]. On the other
19 hand, in support of the thiol-mediated redox reaction model, it was shown that the zebrafish
20 homolog of APE1, which also lacks a Cys65 homolog, does not activate human AP-1 but gains
21 this ability if a Cys is reinstated [115]. For these reasons, and considering recent data on APE1–
22 DNA interactions, an alternative model of transcription factor stimulation has been proposed,
23 discussed in the last section.
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27 Besides transcription factors, APE1 can also stimulate DNA glycosylases in a redox-
28 independent manner. As mentioned above for TDG, many DNA glycosylases bind tightly to
29 their abasic site product. Kinetic characterization of TDG, MBD4, single-strand-specific
30 monofunctional uracil–DNA glycosylase (SMUG1), *N*-methylpurine–DNA glycosylase
31 (MPG/ANPG/AAG), 8-oxoguanine–DNA glycosylase (OGG1), endonuclease III homolog
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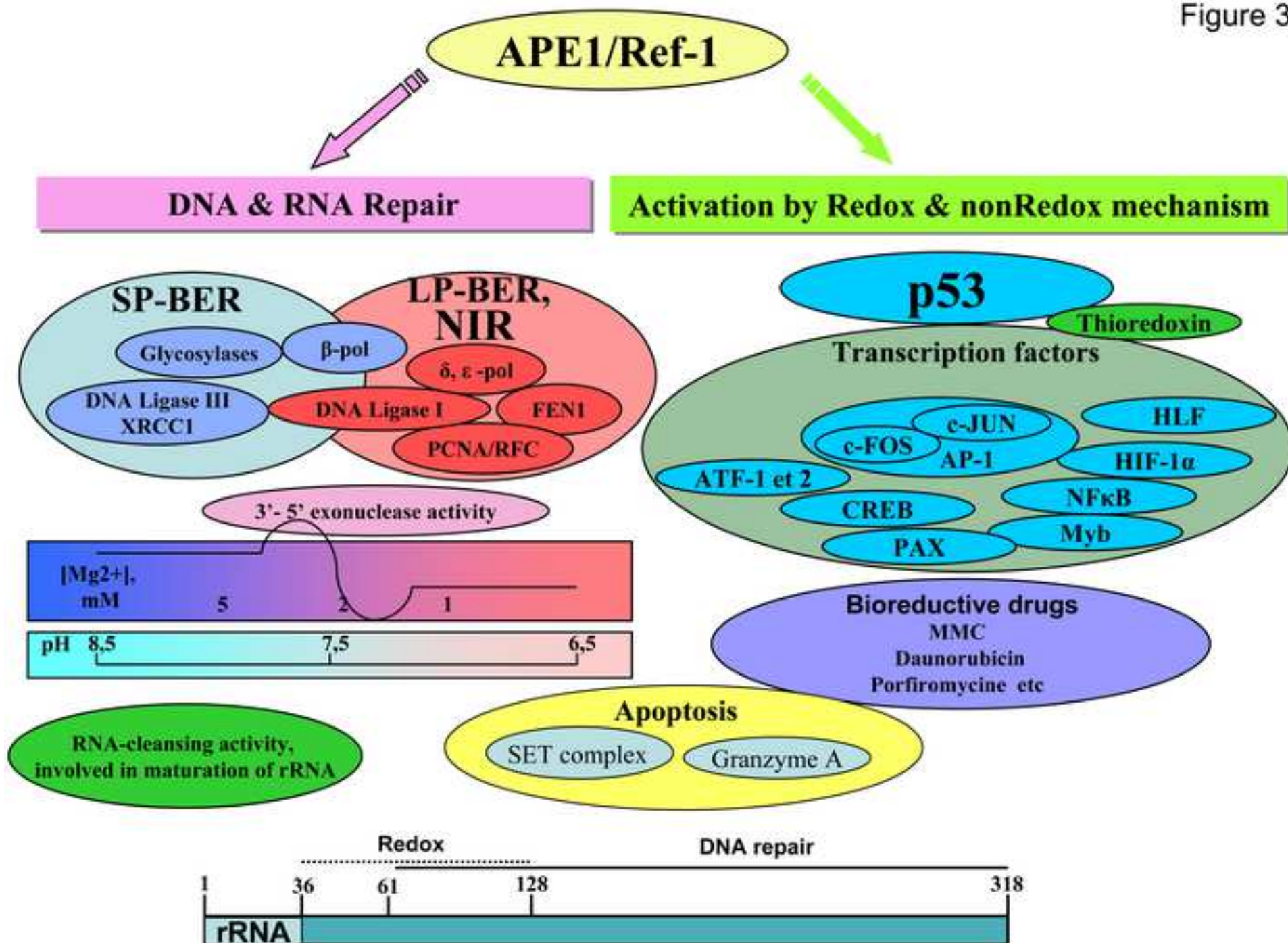
(NTHL1) and MutY homolog (MUTYH) reveals that the product release is rate limiting during the steady-state phase of the reaction [78,143-148]. Full-length APE1 promotes dissociation of the enzyme–product complex, increasing the glycosylase turnover rates [79,148-151]. In contrast, the truncated APE1-NΔ61 mutant lacking the N-tail cannot stimulate the multiple turnover of OGG1, MBD4, and MPG [152].

The N-tail of APE1 is highly enriched in positively charged Lys and Arg residues, which can engage in electrostatic interactions with DNA phosphates, and are involved into multiple posttranslational modifications with a regulatory role. The first 33 residues of the N-tail mediate the binding of APE1 to RNA and to negative Ca²⁺ responsive sequence elements (nCaRE) of certain gene promoters [153-154]. Deletion, mutation, or acetylation of Lys residues within this region increase the AP site cleavage activity, suggesting that the extra charge at the N-tail could retain APE1 on DNA [153,155]. APE1 is acetylated at Lys6 and Lys7 by CBP/p300 and deacetylated by SIRT1, apparently switching between transcription mode (acetylated) and DNA repair mode (deacetylated): the acetylation is required for efficient binding of nCaRE elements [156] and for efficient stimulation of YB-1 and YB1-dependent transcription [138], whereas deacetylated APE1 is bound to XRCC1 [157]. Lys residues 24, 25, and 27 are sites of ubiquitylation, which targets APE1 for degradation and retain it in cytoplasm in a MDM2/p53-dependent manner, possibly indicating shutdown of DNA repair in apoptosis-committed cells [158]. Interestingly, in a specific form of apoptosis initiated by granzyme A protease, APE1 is cleaved at Lys31 and then fully degraded [159]. The first 13 residues of APE1 contain the nuclear localization signal [160]; at the same time, the intact N-tail is suggested to mask the mitochondrial targeting sequence located in the C-terminal region of the polypeptide [161].

In addition to BER, APE1 plays a key role in the removal of damaged DNA bases *via* the nucleotide incision repair (NIR) pathway [162-164] (Figure 3). In BER, APE1 acts downstream of DNA glycosylases by incising a DNA duplex at AP sites and removing 3'-blocking sugar–phosphate moieties. In NIR, APE1 bypasses the glycosylase action and directly generates a

Figure 3
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Figure 3



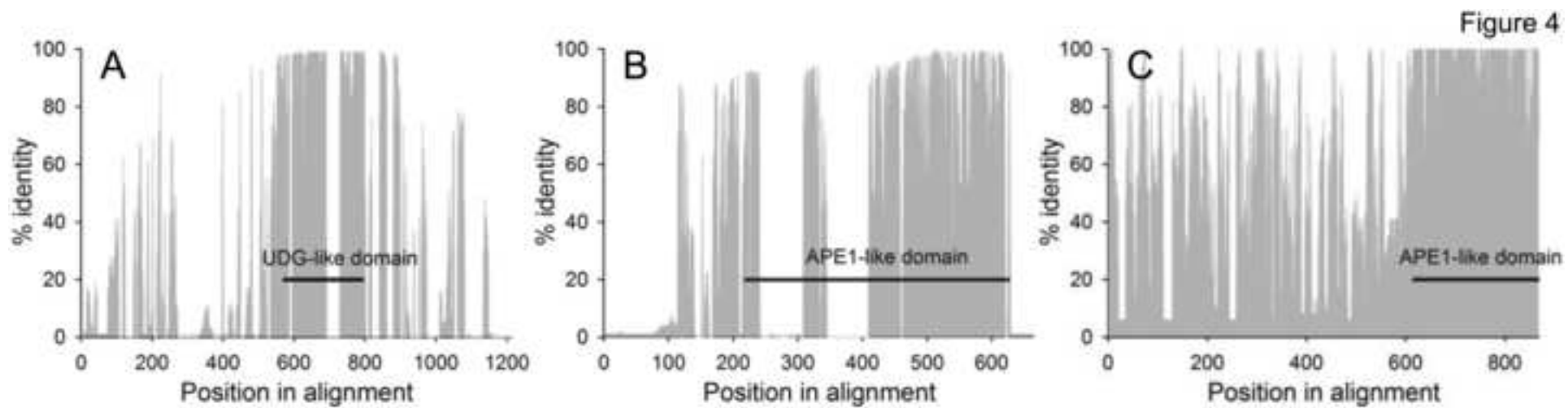
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single-strand break with a 5'-dangling damaged nucleotide and a 3'-hydroxyl group [163,165]. In the NIR mode, APE1 can recognize diverse types of DNA base lesions including α -anomeric deoxynucleotides (α dN), oxidized pyrimidines [163,166], formamidopyrimidines [167], exocyclic DNA bases and uracil [168-169], bulky lesions such as benzene-derived DNA adducts [170] and a UV-induced 6-4 photoproduct [171]. The chemical structures of these DNA lesions have very little in common, implying that in, contrary to DNA glycosylases, APE1 tends to recognize damage-induced structural distortions of the DNA helix and not the modified base itself. The BER/NIR dichotomy in the APE1 action is modulated by concentrations of divalent cations, pH, and ionic strength in an apparently allosteric manner [163]. Changes in intracellular Mg^{2+} concentration induce conformational rearrangements in APE1, changing its DNA substrate specificity [163,172]. Notably, the truncated APE1-N Δ 61 protein exhibits a dramatic decrease in the NIR activity, suggesting a role of the N-tail in its regulation [163,166].

4. Evolution of tails

Given the apparent involvement of TDG and APE1 into transcription regulation and epigenetic demethylation in mammals, and possible involvement of the tails of the mammalian proteins into the interactions important for these functions, it is of great interest to follow the evolution of sequences extending the catalytic domains of these proteins. As the features of cytosine methylation and active demethylation are quite different between vertebrates and other eukaryotes, we have limited the initial analysis to Deuterostomia, a clade that includes vertebrates, echinoderms, and several minor taxa. A comparison of 731 sequences from genomes of 290 Deuterostomia species retrieved from the NCBI RefSeq database as having a TDG, Mug-like catalytic core showed that TDG is present across this clade, and, as expected, possesses a conserved catalytic core, with the similarity in the tails fading towards the polypeptide termini (Figure 4A).

Figure 4
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1 suggested that the “redox function” might be featured only in mammals [115]. We have analyzed
2
3 290 unique sequences containing the APE1-like domain from Deuterostomia (184 species,
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5 including 97 mammals, 81 vertebrate non-mammals, and 6 non-vertebrate chordates and non-
6
7 chordate Deuterostomia). Whereas the catalytic domain was well conserved, the N-tails were
8
9 much more diverse (Figure 4B). Most of this diversity was probably produced by alternative
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11 translation start sites and exon use in different species: whereas all four known RNA isoforms of
12
13 human APE1 encode the same polypeptide, at least 20 various types of N-tails could be
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15 identified in vertebrates. Cys in the position corresponding to human Cys65 was present only in
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17 mammals, with Ser/Thr occupying this position in all analyzed non-mammalian vertebrates.
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22
23 In the absence of clear homology, the basic secondary structure elements can still be
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25 conserved between proteins from different species. To address whether this is the case for APE1
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27 N-tail, we had run secondary structure predictions for 45 vertebrate APE1 sequences (23
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29 mammalian and 22 non-mammalian species). The predicted structures (classified as helix, sheet,
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31 or random coil for each position) were compared using the Levenstein distance, a string metric
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33 for measuring the difference between non-numerical sequences. Moderate conservation of the N-
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35 tail secondary structure for mammals was revealed, while lacking in other vertebrates (Figure
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37 5A). In comparison, the structures of C-terminal sequences of equal lengths are strongly
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39 conserved in the clusters of mammals (with an exception of marsupials) and non-mammalian
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41 vertebrates (Figure 5B). Overall, it seems that N-tails in mammalian APE1 proteins have indeed
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43 evolved some functions making them distinct from their non-mammalian counterparts.
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50 Of known AP endonucleases, at least one more has a long functionally significant
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52 extension: the *Drosophila* protein, Rrp1, carries a BER function in its C-terminal part and a
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54 homologous recombination function (albeit of unknown mechanism) in its N-terminal part. Rrp1
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56 was first isolated from fruit fly embryos as a protein capable of promoting strand annealing and
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58 strand exchange between single- and double-stranded DNA molecules in an energy-independent
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Figure 5
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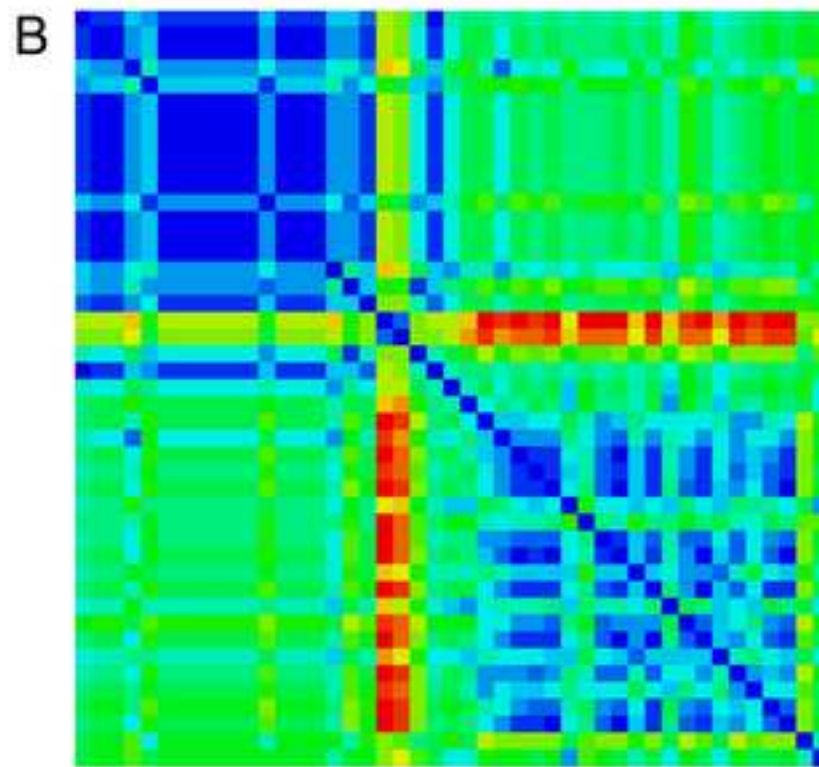
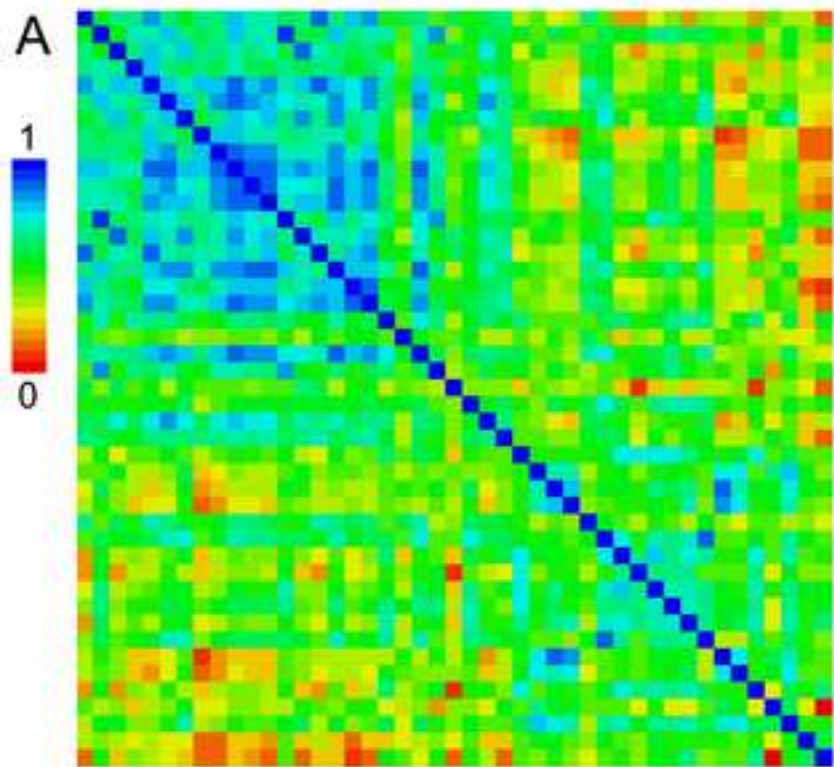


Figure 5

1 manner [173]. The compact C-terminal part (251 aa long; [Figure 1B](#)) was later shown to bear
2 homology with EEP superfamily AP endonucleases [174]. It catalyzes cleavage of AP sites,
3 resection of blunt or recessed 3'-ends in a DNA duplex, possesses 3'-phosphatase and 3'-
4 phosphodiesterase activities on recessed damaged 3'-ends [175-178] and complements the
5 oxidation- and alkylation-sensitive phenotype of AP endonuclease-null *E. coli* [179], showing all
6 attributes of a functional AP endonuclease participating in BER. Although no structural
7 information on Rrp1 is available, limited proteolysis combined with circular dichroism and
8 sedimentation velocity analysis indicate that the N-tail (426 aa) is mostly unstructured [180]. The
9 physiological role of Rrp1 in recombination is not clear: overexpression of full-length Rrp1 in
10 *Drosophila* larvae reduces rather than increases the level of DNA damage-induced genetic
11 marker conversion but this could reflect more efficient repair by BER and not suppression of
12 recombination [181]. Rrp1 might contribute to recombination indirectly: its N-tail interacts with
13 both Rev3 and Rev7 subunits of DNA polymerase ζ [182], an error-prone DNA polymerase that
14 extends from terminally mismatched DNA primers and, from genetic evidence, is involved in
15 homologous recombination in *Drosophila* [183].
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35 A protein sequence database search reveals that functional AP endonucleases are linked
36 with the recombination functional part over a very narrow phylogenetic range. Rrp1 sequences
37 are available for 25 *Drosophila* species, with up to four protein isoforms predicted for some
38 genomes. Their comparison clearly shows that the N-tail is much less conserved than the C-
39 terminal APE1 homology domain ([Figure 4C](#)). Thus, even within the same genus, the N-terminal
40 sequences, while still recognized as homologous, diverged much faster than AP endo-like
41 domain sequences. Other Diptera also possess APE1-like AP endonucleases with long N-tails,
42 their homology to the recombination part of *Drosophila* proteins are below the limit of positive
43 identification. In other insects, the length of the N-tail of Rrp1/APE1 fluctuate greatly. In
44 Lepidoptera and Coleoptera, the Rrp1 part was as extensive or even longer as in Diptera, while
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1 in other insect orders it could be less than 70 aa. Clearly, the recombination function cannot be
2 universally ascribed to the N-tail of insect APE1 proteins.

3
4 The failure to obtain crystal structures of TDG and APE1 tails suggests that they are
5 likely disordered. We have used ESpritz bi-directional recursive neural network [184] trained on
6 the training set of the DisProt database of partially or completely disordered proteins [185] to
7 predict disorder in the tails of human APE1 and TDG and *D. melanogaster* Rrp1. Indeed, the
8 tails showed higher disorder probability than the catalytic cores, and the boundaries of the
9 predicted disordered regions well coincided with the borders between the core and the tails
10 (Figure 6). It is known that terminal regions of protein-coding sequences in both mammals and
11 insects are evolutionarily more labile, accumulating more non-synonymous mutations than the
12 central part that presumably contains the functional core of the protein [186]. For an important
13 class of proteins – those composed a mammalian cell mRNA interactome – high intrinsic
14 disorder was shown to be associated with the enrichment in short repetitive amino acid motifs
15 [187]. To assess the representation of short repeats in the tails and the core of APE1 and related
16 AP endonucleases we have analyzed 2766 sequences containing the APE1-like domain using
17 rapid automatic detection and alignment of repeats (RADAR) algorithm [188]. Of these, 370
18 contained an N-tail > 40 aa long (only 22 sequences contained a C-tail, which was not enough
19 for a valid comparison). As an example of such analysis, Figure 7A and B illustrates the results
20 for Rrp1, in which 74% of the N-tail turned out to be covered by short repeats. N-tails were
21 significantly enriched in short repeats in comparison with the catalytic cores ($p < 10^{-6}$,
22 Kolmogorov–Smirnov test; Figure 7C). TDG sequences produced a similar picture for both N-
23 and C-tails. Thus, as in the case of mRNA-binding proteins, disorder in the tail of APE1-like
24 endonucleases could be due to the presence of repetitive protein motifs. It should be emphasized
25 that the disorder by itself may have functional significance and the evolution might favor
26 accumulation of short diverged repeats in the appropriate protein regions.
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Figure 6
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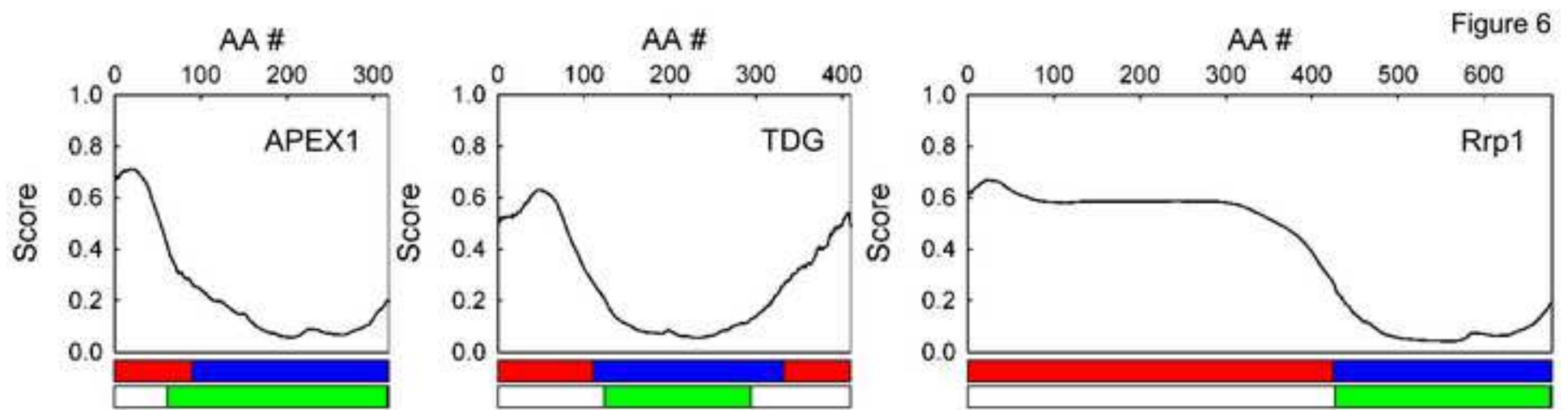


Figure 7
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Figure 7

A

No. of Repeats	Total Score	Length	Diagonal	BW-From	BW-To	Level
2	56.67	21	23	344	366	1

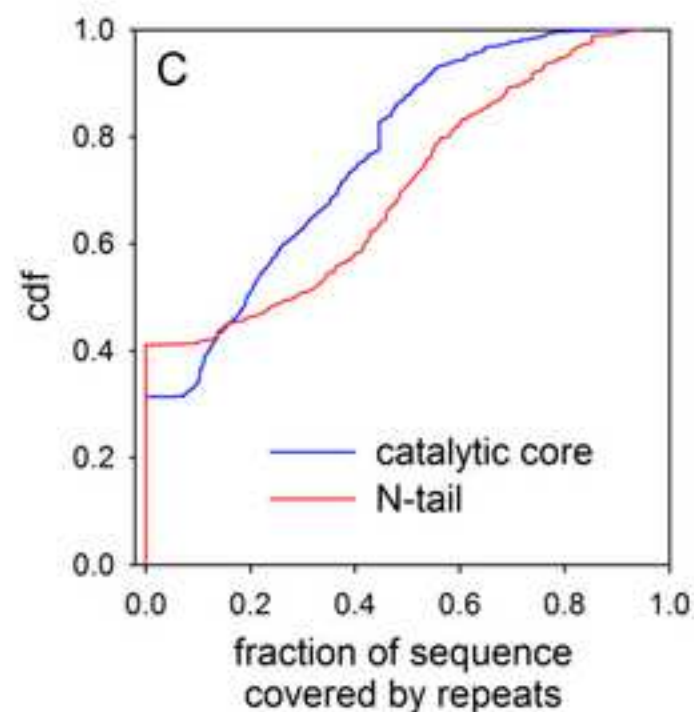
333- 359 (31.08/ 7.64) **GV..VEE**eakp**stETKPAKGRKKAPVKA**E
 360- 385 (25.58/ 8.31) **DVedIEE...**aa**eESKPARGRKKAAAKAE**

No. of Repeats	Total Score	Length	Diagonal	BW-From	BW-To	Level
5	260.65	45	150	144	188	5

3- 44 (42.35/10.98) ...**RVKAVK.KQAEALASEPTDP**...**TPNANGnG**...**VDE**.....**NADSAAEELKVP**
 45- 94 (51.84/15.60) **AKGKPRARKATKTAVSAENSEEVE**pqka**PTAAAR.G**...**KKK**.....**QPKDTE**NGQ**ME**
 144- 188 (71.53/25.19) **PKGRAKAEKPTNAQAKGRKRKEL**...**PAEANG.G**...**AEE**.....**AAEPPKQ**AR**KE**
 191- 234 (53.30/16.31) **PTLKEQAE**PG**TISKEKVQK.AET**...**AAKRAR.G**...**TKR**.....**LADSEIAAALDE**
 235- 314 (41.63/10.63) **PEVDEVPPKAASKRAKKGK**M**VEP**...**SPETVG**.Dfqs**vQEE**vesppktaaap**kk**ra**kk**tt**ng**etave**lep**kt**KAKPTKQ**RA**KE**

B

MPRVKAVKKQAEALASEPTDPTPNANGNGVDENADSAAEELKVP**AKGKPR** 50
ARKATKTAVSAENSEEVEPQKAPTAAARGKKK**QPKDTE**NGQ**MEV**VAKPK 100
 GRAKKATAEAEPEPKVDLPAGKATK**PRAKKEPT**PAPDEV**TSSPPK**GRAKA 150
EKPTNAQAKGRKRKELPAEANGGAEEAAEPPK**Q**AR**KEAVPTLKEQAE**PG 200
TISKEKVQKAETAAKRARGTKRLADSEIAAALDE**PEVDEVPPKAASKRAK** 250
 KGKMVEPS**PETVGD**FQSVQ**EEVES**PPK**TAAAPK**KRAK**TTNG**ETAVELEP 300
 KTKAKPT**KQRAKKEG**KEPAPGKKQKKSADKENG**VVEE**EAK**PSTETKPAK**G 350
RKKAPVKAEDVEDIEEAAEESKPARGRKKAAAKAE**EPDVDEESGSK**TT**TK** 400
 AKKAETKTT**VTLDKDAFAL**PADKEFN**LKIC**SWNVAGLRAWL**KKDGLQLID** 450
LEEPDIFCLQETKCANDQLPEEV**TRLPGYHPY**WLCMPGGYAGVAI**YSKIM** 500
 PIHVEY**GIGNEEFDDVGR**MITAEY**EKFYLIN**VYVPNSGRKLVNLE**PRMRW** 550
EKLFQAYVKKLDAL**KPVVICGDMNV**SHMPIDLEN**PKNNTKNAGFTQ**EE**RD** 600
KMTELLGLGFVD**TFRHLYPDR**KGAY**TFW**TYMANARAR**NVGWR**LDYCLVSE 650
RFVPKVVEHE**IRSQCLGSDHC**PI**TIF**ENI 679



5. APE1 and TDG: cooperative mediators?

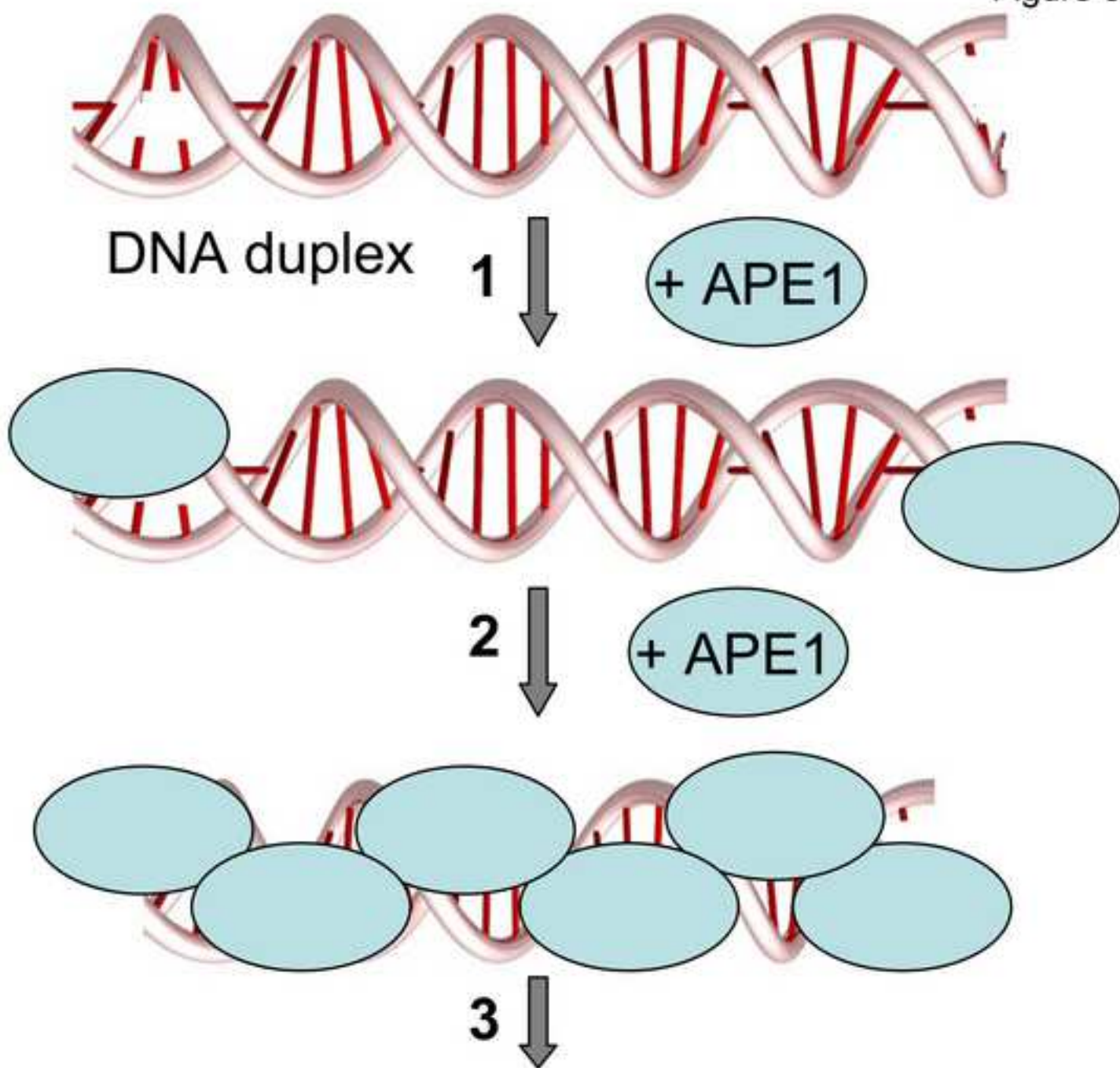
1 As discussed above, the N-tail of APE1 is indispensable for the functions that appear very
2 different mechanistically, namely the stimulation of DNA binding by transcription factors,
3 turnover of DNA glycosylases, and the BER/NIR switch. Recent observations brought forward a
4 somewhat unexpected hypothesis that unifies all these functions under a model of APE1
5 multimerization on DNA. Transmission electron microscopy and atomic force microscopy
6 revealed that the full-length APE1 can form filaments on both undamaged and damaged DNA
7 fragments and introduce bends and kinks that can be used to detect an AP site [149,189]. Several
8 APE1 molecules was observed to bind both undamaged and damaged oligonucleotides in an
9 apparently cooperative mode, with single-stranded DNA or nicks acting as preferential binding
10 sites [189]. Combined with the quantitative proteomics estimates revealing that APE1 is an
11 abundant protein in human cells, exceeding the amount of most DNA glycosylases and
12 transcription factors by > 2 orders of magnitude [190-191], these observations suggest that APE1
13 monomers could dynamically assemble on DNA, starting from DNA nicks, gaps or breaks.
14 Random binding and dissociation of terminal APE1 monomers effectively amounts to a
15 stochastic walk of the multimer along the DNA contour, mechanistically equivalent to DNA
16 scanning by individual molecules of many DNA repair proteins, restriction endonucleases and
17 transcription factors [192-193]. In this mode, an APE1 multimer may “search” for some specific
18 sites in DNA. Importantly, truncated APE1-N Δ 61 multimerizes on DNA much less efficiently
19 than the full-length protein [152], and this decreased capacity strongly correlates with the
20 reduced ability of APE1-N Δ 61 to perform NIR and to stimulate transcription factors and DNA
21 glycosylases [110-111,126,152,163,189].

22 In addition, the modeled structure of the full-length APE1 could be docked into
23 oligomers with translational symmetry, in which the DNA-binding site is exposed, consistent
24 with the APE1 multimer assembly on DNA [189]. Binding of proteins in the DNA minor groove
25 is known to induce groove dehydration and easy DNA bending [194]. Based on these

1 observations, it has been proposed that APE1 multimerization induces DNA helix distortion that
2 exacerbates pre-existing anisotropic flexibility at AP sites, oxidized nucleotides and intrinsically
3 bent DNA, which in turn enables NIR activity, stimulation of DNA binding by transcription
4 factors, and stimulation of DNA glycosylases turnover [152,189] (Figure 8).
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8 What could be the mechanism responsible for the stimulation of transcription factors by
9 an APE1 multimer? DNA bending due to bound proteins also induces distortions at neighboring
10 sequences, which may promote the binding of transcription factors; a good example is p53
11 binding to its cognate sequence, which is favored by the DNA curvature induced by HMGB1
12 chromatin architectural protein [195]. Generally, the proteins that induce DNA bending exhibit
13 better affinity when their recognition sites are located in intrinsically bent or flexible regions, and
14 intrinsic DNA bending plays important role in transcription regulation [196-198]. Both NF- κ B
15 and AP-1 have been demonstrated to bend DNA [199-200]. Even though the DNA bending by
16 isolated AP-1 was debated [201], accessory factors such as NFAT1 assist bending and AP-1
17 binding to its recognition sites [202-203]. Essentially, bending induced by APE1 can play the
18 same role, sculpting DNA for easier binding of transcription factors. Although no comprehensive
19 analysis of sequence preferences was done, APE1 may itself favor intrinsically bent DNA: it
20 binds to nCaRE-A and nCaRE-B sites in the promoters of the parathyroid hormone (*PTH*) gene
21 and the own *APE1* gene [154,204-205]. nCaRE sequences contain intrinsically bent A-tracts
22 d(TTTTTG)/d(CAAAAA) flanking the recognition site for nCaRE-binding proteins. Overall,
23 APE1 might specifically bind to and multimerize on intrinsically bent cognate sites of
24 transcription factors and promote further DNA distortion, which in turn stimulates transcription
25 factor binding.
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51 There are also some evidence for functionally relevant oligomerization of DNA
52 glycosylases, including TDG. Structural studies revealed that the catalytic domain of TDG binds
53 to 23–28-mer DNA duplexes in a 2:1 ratio, with a second TDG molecule bound in a non-
54 catalytic orientation, whereas the protein dimer complex cannot be formed on a short 15-mer
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Stimulation of enzymatic turnover of DNA glycosylases and DNA binding of transcription factors.

1 DNA duplex [55,72]. Full-length TDG forms dimers on short duplexes with much higher
2 efficiency, suggesting that N- and C-tails participate in the assembly of TDG on DNA. In
3 relation to these observations, it has been shown that *E. coli* Mug, human O⁶-alkylguanine-DNA
4 alkyltransferase (MGMT), and N-truncated *E. coli* MutY DNA glycosylase exhibit cooperative
5 binding to DNA substrates and can form protein oligomers on DNA with 4–12 nucleotides
6 periodicity [206-208]. It was hypothesized that the cooperative mode of binding could enable
7 more efficient lesion search or protect repair intermediates before handing them over to
8 downstream BER machinery. The stoichiometric difference between TDG complexes formed on
9 on 28- and 15-mer DNA duplexes has no measurable effect on the U excision activity [72].
10 However, TDG does not excise T opposite ethenoadenine or hypoxanthine, or 8-oxoadenine
11 opposite to T from 15-mer duplexes, suggesting that the cooperative binding of TDG to DNA
12 may be required for efficient processing of a wider range of DNA substrates [103]. At present,
13 the molecular mechanism and structural basis for this molecular recognition phenomenon remain
14 unclear.
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32 At least for APE1, the ability to assemble on DNA can have profound consequences for
33 gene regulation. A testable prediction from this model is that regulatory sites exhibiting
34 anisotropic flexibility or intrinsic DNA bending would be prone to APE1-mediated stimulation
35 of DNA binding by transcription factors. Similarly, multimerization of APE1 at specific
36 regulatory elements may enhance the removal of oxidized bases via stimulation of the DNA
37 glycosylases. Noteworthy, it has been shown that activation of several promoters, including
38 those responsive to ER α and c-Myc and also those containing potential quadruplex-forming
39 sequences, requires guanine oxidation followed by BER in an OGG1, APE1-dependent manner
40 [209-211]. Conversely, displacement of OGG1 by APE1 could compete with the activation of a
41 number of immune- and stress-responsive genes, that relies on reversibly oxidized OGG1 that
42 binds but does not excise 8-oxoguanine [212-213]. Of course, stimulation of TDG and MBD4 by
43 APE1 may also contribute to gene regulation by TET-dependent active demethylation, both
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1 through the erasure of mC marks and through recruitment of poly(ADP-ribose) polymerase 1
2 (PARP1) to the nicked repair intermediates, which poly(ADP-ribose)lates histones and recruits
3 of chromatin remodeling factors. Finally, DNA curvature introduced by APE1 and enhanced by
4 transcription factors binding, or PARP1 recruited to the APE1-produced nicks could promote
5 DNA loop formation and interactions with distantly located enhancer or insulator elements.
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10 11 12 13 **6. Conclusion** 14

15
16 The possibility of BER proteins combining their BER function with some other activity
17 and containing structurally separated determinants for both, is attractive in the light of recent
18 discoveries of the role of the BER pathway in processes not involved with genome integrity
19 maintenance. In particular, in the last years it has been firmly established that mammalian cells
20 rely on TDG-initiated BER to actively demethylate epigenetically regulated genes. Mammalian
21 zygotes or reprogrammed cells undergo massive demethylation through this mechanism to erase
22 epigenetic marks and re-establish pluripotency. Another well-documented regulatory role
23 belongs to APE1, which, in addition to its role in BER, stimulates DNA binding by many
24 transcription factors.
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38 Both TDG and APE1 have homologs in bacteria but have a notable feature in the
39 presence of extended tails in addition to the catalytic cores. Notably, their archaeal homologs
40 also lack the tails, indicating that these structural elements emerged during the evolution of
41 eukaryotes. However, even in eukaryotes, the tails of TDG and APE1 homologs are much more
42 diverse than their catalytic cores, as evidenced, for example, by human APE1 and *Drosophila*
43 Rrp1 that have totally different sequences, structures, and functions ascribed to their N-tails. The
44 evolution of the tails might proceed through accumulation and fast divergence of short repeats,
45 which would be initially close to neutral outside of the catalytic cores, and then switch to a
46 directional selection mode once some useful function was acquired.
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1 It is clear that tails of TDG and APE1 are involved in a multitude of protein–DNA and
2 protein–protein interactions and serve as elements for regulatory protein modification. Currently,
3 the mechanistic understanding of the functions of TDG and APE1 tails is limited by the lack of
4 structural information. Given the intrinsically disordered nature of the tails, their structure in
5 isolation is unlikely to be determined. However, it is possible that the conformations of the tails
6 are better defined in some stable or metastable protein–protein complexes, including those
7 involved in epigenetic DNA demethylation, which can be analyzed by conventional methods of
8 structural biology. Moreover, new experimental methods to study mechanistically important
9 features of intrinsically disordered protein regions are being developed using tools from NMR
10 [214], single-molecule fluorescence microscopy [215], isotope exchange mass spectrometry
11 [216], etc. The characterization of the tails' behavior is especially important in view of the
12 explosive development of a new paradigm of biomolecular ensemble organization based on
13 liquid phase separation, which is now believed to underlie many cell regulation processes
14 including DNA repair and chromatin remodeling [217-220]. It cannot be excluded that tail-
15 dependent assembly of APE1 and TDG aggregates on DNA might be involved in the liquid
16 demixing in DNA repair and active epigenetic demethylation.
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48 (PRC CNRS/RFBR n1074 REDOBER).
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Legends to figures

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3 **Figure 1.** Schematic domain structure of *E. coli* Mug and human TDG (A) and *E. coli*
4 Xth, human APE1, and *Drosophila* Rrp1 (B). The shaded blocks represent the enzymes'
5 conserved catalytic domains. The lines below indicate structurally characterized part of the
6 human proteins. Functionally important Lys330 in TDG and Cys65 in APE1 are marked by
7 arrows.
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13 **Figure 2.** Schematic representation of the active DNA demethylation pathways. mC can
14 be hydroxylated by TET1, TET2, or TET3 proteins to form hmC or further oxidized fC and caC.
15 In addition, mC and hmC can be deaminated by the AID/APOBEC family members to form T
16 and hmU. Removal of these demethylation intermediates by DNA glycosylases in the BER
17 pathway results in the erasure of mC and its replacement with C. Alternatively, de novo DNA
18 methyltransferases DNMT3a/3b are able to directly convert hmC to C without base excision and
19 repair synthesis. The inset shows the essential short-patch BER pathway in mammalian cells: the
20 modified base (red) is removed by a DNA glycosylase, an AP endonuclease then nicks 5' to the
21 nascent AP site, DNA polymerase β inserts the correct nucleotide (green) and excises the
22 remaining deoxyribose phosphate, and DNA ligase III α seals the nick.
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41 **Figure 3.** Mammalian AP endonuclease 1 (APE1), an essential multifunctional repair
42 protein. (Left) APE1 is involved in the BER and NIR pathways to remove non-bulky DNA base
43 damage and degrades damaged abasic RNA. BER/NIR switch and DNA substrate specificity of
44 APE1 are modulated by concentrations of divalent cations, pH and ionic strength. (Right) APE1
45 stimulates DNA binding of various transcription factors *via* redox- and nonredox-dependent
46 mechanisms and also participates in the activation of bioreductive drugs. APE1 is a member of
47 the SET complex and promotes the Granzyme A activated cell death response. (Bottom)
48 Schematic structure of APE1 showing amino acids involved in nuclear localisation signal (NLS),
49 redox function and DNA repair activities.
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1 **Figure 4.** Conservation plot of the alignment of TDG (A, 731 sequences from genomes
2 of 290 Deuterostomia species), APE1 (B, 290 sequences from 184 Deuterostomia species) and
3 Rrp1 (C, 49 sequences from genomes of 25 *Drosophila* species). The alignment was done using
4 MAFFT [221]. The fraction of the most frequent residue in every position is shown.
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8 **Figure 5.** Conservation of the secondary structures at the N- and C-terminus of APE1
9 sequences. Secondary structure predictions were run for 45 vertebrate APE1 sequences (23
10 mammalian and 22 non-mammalian species, limited to one per taxonomic order to minimize
11 similarity by descent) that have the N-tail > 40 aa long (44–67 in the set) using GOR IV [222].
12 The predicted strings of secondary structure assignments were compared by Levenstein distance
13 between them [223]. Color-coded normalized distance matrices are shown for the complete N-
14 tails (A); and for the last 70 C-terminal residues in the catalytic domain (B).
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25 **Figure 6.** Predicted disordered regions in human APE1 and TDG and *D. melanogaster*
26 Rrp1. ESpritz probability score is plotted over the length of the protein. Best weighted score
27 method for disorder prediction was used [224]. Red bars show the predicted disordered regions,
28 blue bars, the ordered regions. Green bars show the positions of the catalytic cores.
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35 **Figure 7.** Detection of repeats in the APE1 family N-tails. (A), list of repeats in
36 *Drosophila* Rrp1 identified by RADAR [188]. (B), repeats mapped onto the Rrp1 sequence (red,
37 catalytic domain; cyan and magenta, alternating level 1 repeats; yellow and green, alternating
38 level 5 repeats). (C), cumulative distribution function (cdf) for the fraction of the length of the
39 catalytic domain and the N-tail covered by repeats in 2766 APE1 proteins (370 containing N-
40 tails longer than 40 aa).
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49 **Figure 8.** Proposed unified mechanism of the APE1-catalyzed stimulation of DNA
50 glycosylases and transcription factors. (1) APE1 binds to transient single-stranded structures
51 present at the termini of duplex DNA. (2) Multiple molecules of APE1 assemble on the DNA
52 fragment and form multimer complexes. (3) APE1 multimers induce conformational changes in
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the DNA helix, which in turn stimulates DNA glycosylase activities and sequence-specific binding of transcription factors to DNA.

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